Claims

1. A compound of formula (I):

(I)

wherein:

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 \mathbb{R}^1 is halo, cyano, \mathbb{C}_{1-3} alkyl or \mathbb{C}_{1-3} alkoxy;

p is 0-2; wherein the values of R¹ may be the same or different;

R² is hydrogen, C₁₋₄alkyl, C₂₋₄alkenyl, C₂₋₄alkynyl, C₃₋₆cycloalkyl,

C₃₋₆cycloalkylC₁₋₃alkyl, a heterocyclyl or heterocyclylC₁₋₃alkyl; wherein R² may be optionally substituted on carbon by one or more methyl, ethyl, methoxy, ethoxy, propoxy, trifluoromethyl, trifluoromethoxy, 2,2,2-trifluoroethoxy or cyclopropylmethoxy; and wherein if said heterocyclyl contains an -NH- moiety that nitrogen may be optionally substituted by one or more methyl, ethyl, acetyl, 2,2,2-trifluoroethyl or methoxyethyl:

R³ is hydrogen, halo or cyano;

 \mathbf{R}^4 is C_{1-6} alkyl or C_{1-6} alkoxy C_{1-6} alkyl;

 ${\bf R}^5$ is C_{1-6} alkyl or C_{2-6} alkenyl; wherein ${\bf R}^5$ may be optionally substituted on carbon by one or more methoxy, ethoxy, propoxy, trifluoromethyl, trifluoromethoxy,

2,2,2-trifluoroethoxy or cyclopropylmethoxy;

 \mathbb{R}^6 is halo or \mathbb{C}_{1-4} alkyl;

or a pharmaceutically acceptable salt or an *in vivo* hydrolysable ester thereof; with the proviso that if R^4 , R^5 and R^6 are all methyl then R^2 is not hydrogen, optionally substituted C_{1-4} alkyl or C_{3-6} cycloalkyl.

25 2. A compound of formula (I) according to claim 1 wherein p is 0; or a pharmaceutically acceptable salt or an *in vivo* hydrolysable ester thereof.

3. A compound of formula (I) according to claim 1 or 2 wherein R^2 is hydrogen, C_{1-4} alkyl or heterocyclyl C_{1-3} alkyl; wherein R^2 may be optionally substituted on carbon by one or more methoxy or ethoxy; or a pharmaceutically acceptable salt or an *in vivo* hydrolysable ester thereof.

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- 4. A compound of formula (I) according to anyone of claims 1-3 wherein R³ is hydrogen or halo; or a pharmaceutically acceptable salt or an *in vivo* hydrolysable ester thereof.
- 5. A compound of formula (I) according to anyone of claims 1-4 wherein R⁴ is C₁₋₄alkyl; or a pharmaceutically acceptable salt or an *in vivo* hydrolysable ester thereof.
 - 6. A compound of formula (I) according to anyone of claims 1-5 wherein R^5 is C_{1-6} alkyl; or a pharmaceutically acceptable salt or an *in vivo* hydrolysable ester thereof.
- 7. A compound of formula (I) according to anyone of claims 1-6 wherein R⁶ is methyl or halo; or a pharmaceutically acceptable salt or an *in vivo* hydrolysable ester thereof.
 - 8. A compound of formula (I) (as depicted in claim 1) wherein: p is 0;
- 20 R² is hydrogen, 2-methoxyethyl, methyl, 3-methoxypropyl or 2-ethoxyethyl or 2-pyrazol-1-ylethyl;

R³ is hydrogen or bromo;

R⁴ is methyl, ethyl or isopropyl;

R⁵ is methyl or ethyl;

25 R⁶ is methyl or bromo;

or a pharmaceutically acceptable salt or an *in vivo* hydrolysable ester thereof; with the proviso that if R^4 , R^5 and R^6 are all methyl then R^2 is not hydrogen, 2-methoxyethyl, methyl, 3-methoxypropyl or 2-ethoxyethyl.

9. A compound of formula (I) (as depicted in claim 1) selected from:
 4-(1,2-diethyl-4-methylimidazol-5-yl)-2-{4-[N-(2-methoxyethyl)sulphamoyl]anilino}
 pyrimidine;

- 4-(1-ethyl-2,4-dimethylimidazol-5-yl)-2-{4-[*N*-(2-ethoxyethyl)sulphamoyl]anilino} pyrimidine;
- 4-(1-ethyl-2,4-dimethylimidazol-5-yl)-2-{4-[*N*-(3-methoxypropyl)sulphamoyl]anilino} pyrimidine;
- 5 4-(1-ethyl-2,4-dimethylimidazol-5-yl)-2-{4-[*N*-(2-methoxyethyl)sulphamoyl]anilino} pyrimidine;
 - 4-(1-ethyl-2,4-dimethylimidazol-5-yl)-2-{4-[N-(methyl)sulphamoyl]anilino}pyrimidine;
 - 4-(1-ethyl-2,4-dimethylimidazol-5-yl)-2-(4-sulphamoylanilino)pyrimidine;
 - $4-(1-isopropyl-2,4-dimethylimidazol-5-yl)-2-\{4-[N-(2-methoxyethyl)sulphamoyl] anilino\}$
- 10 pyrimidine;
 - 4-(1-isopropyl-2,4-dimethylimidazol-5-yl)-2-{4-[*N*-(3-methoxypropyl)sulphamoyl]anilino} pyrimidine;
 - 4-(1-isopropyl-2,4-dimethylimidazol-5-yl)-2-(4-sulphamoylanilino)pyrimidine;
 - 4-(1-isopropyl-2,4-dimethylimidazol-5-yl)-2-{4-[N-(2-ethoxyethyl)sulphamoyl]anilino}
- 15 pyrimidine;
 - 4-(1-isopropyl-2,4-dimethylimidazol-5-yl)-2-{4-[N-(methyl)sulphamoyl]anilino}pyrimidine;
 - 4-(1,2,4-trimethylimidazol-5-yl)-2-{4-[*N*-(2-pyrazol-1-ylethyl)sulphamoyl]anilino} pyrimidine; or
 - $4-(4-bromo-1,2-dimethylimidazol-5-yl)-2-\{4-[\textit{N-}(2-methoxyethyl)sulphamoyl] anilino \} \\$
- 20 pyrimidine;
 - or a pharmaceutically acceptable salt or an in vivo hydrolysable ester thereof.
- 10. A process for preparing a compound of formula (I) or a pharmaceutically acceptable salt or an *in vivo* hydrolysable ester thereof which process (wherein R¹, R², R³, R⁴, R⁵, R⁶ and p are, unless otherwise specified, as defined in claim 1) comprises of:
 Process a) reaction of a pyrimidine of formula (II):

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$$\begin{array}{ccccc}
R^{3} & & & & & \\
R^{4} & & & & & \\
R^{5} & & & & & \\
\end{array}$$
(II)

wherein L is a displaceable group; with an aniline of formula (III):

$$(III)$$

$$(R^1)_p$$

$$H$$

$$O$$

$$O$$

$$R^2$$

or

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Process b) reacting a compound of formula (IV):

$$\begin{array}{c|c} H \\ N \\ N \\ N \\ \end{array}$$

$$\begin{array}{c} H \\ N \\ \end{array}$$

$$\begin{array}{c} (R^1)_p \\ \\ N \\ \end{array}$$

$$\begin{array}{c} H \\ N \\ \end{array}$$

$$\begin{array}{c} R^2 \\ \end{array}$$

$$\begin{array}{c} (IV) \\ \end{array}$$

with a compound of formula (V):

$$R^3$$
 R^4
 R^5
 R^5
 R^8
 R^8
 R^8
 R^8
 R^8
 R^8

wherein T is O or S; R^{x} may be the same or different and is C_{1-6} alkyl;

Process c) reacting a pyrimidine of formula (VI):

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(VI)

wherein X is a displaceable group; with an amine of formula (VII):

R²-NH₂

(VII)

or

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Process d) reacting a pyrimidine of formula (VIII)

(VIII)

with a compound of formula (IX):

where Y is a displaceable group;

and thereafter if necessary:

- i) converting a compound of the formula (I) into another compound of the formula (I);
 - ii) removing any protecting groups;
 - iii) forming a pharmaceutically acceptable salt or in vivo hydrolysable ester.

- 11. A pharmaceutical composition which comprises a compound of the formula (I), or a pharmaceutically acceptable salt or *in vivo* hydrolysable ester thereof, according to any one of claims 1-9, in association with a pharmaceutically-acceptable diluent or carrier.
- 5 12. A compound of the formula (I), or a pharmaceutically acceptable salt or *in vivo* hydrolysable ester thereof, according to any one of claims 1-9, for use in a method of treatment of the human or animal body by therapy.

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- 13. A compound of the formula (I), or a pharmaceutically acceptable salt or *in vivo* hydrolysable ester thereof, according to any one of claims 1-9, for use as a medicament.
 - 14. The use of a compound of the formula (I), or a pharmaceutically acceptable salt or *in vivo* hydrolysable ester thereof, according to any one of claims 1-9, in the manufacture of a medicament for use in the production of a cell cycle inhibitory (anti-cell-proliferation) effect in a warm-blooded animal such as man.
 - 15. The use of a compound of the formula (I), or a pharmaceutically acceptable salt or *in vivo* hydrolysable ester thereof, according to any one of claims 1-9, in the manufacture of a medicament for use in the treatment of cancers (solid tumours and leukaemias),
- fibroproliferative and differentiative disorders, psoriasis, rheumatoid arthritis, Kaposi's sarcoma, haemangioma, acute and chronic nephropathies, atheroma, atherosclerosis, arterial restenosis, autoimmune diseases, acute and chronic inflammation, bone diseases and ocular diseases with retinal vessel proliferation.
- 25 16. The use of a compound of the formula (I), or a pharmaceutically acceptable salt or *in vivo* hydrolysable ester thereof, according to any one of claims 1-9, in the manufacture of a medicament for use in the treatment of cancer.
- 17. The use according to claim 16 wherein the cancer is selected from leukaemia, breast cancer, lung cancer, colorectal cancer, stomach cancer, prostate cancer, bladder cancer, pancreatic cancer, ovarian cancer, liver cancer, kidney cancer, skin cancer and cancer of the vulva.

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- 18. A compound of the formula (I), or a pharmaceutically acceptable salt or *in vivo* hydrolysable ester thereof, according to any one of claims 1-9, for use in the production of a cell cycle inhibitory (anti-cell-proliferation) effect in a warm-blooded animal such as man.
- 5 19. A compound of the formula (I), or a pharmaceutically acceptable salt or *in vivo* hydrolysable ester thereof, according to any one of claims 1-9, for use in the treatment of cancers (solid tumours and leukaemias), fibroproliferative and differentiative disorders, psoriasis, rheumatoid arthritis, Kaposi's sarcoma, haemangioma, acute and chronic nephropathies, atheroma, atherosclerosis, arterial restenosis, autoimmune diseases, acute and chronic inflammation, bone diseases and ocular diseases with retinal vessel proliferation.
 - 20. A compound of the formula (I), or a pharmaceutically acceptable salt or *in vivo* hydrolysable ester thereof, according to any one of claims 1-9, for use in the treatment of cancer.
 - 21. A compound of the formula (I), or a pharmaceutically acceptable salt or *in vivo* hydrolysable ester thereof, according to any one of claims 1-9, for use in the treatment of leukaemia, breast cancer, lung cancer, colorectal cancer, stomach cancer, prostate cancer, bladder cancer, pancreatic cancer, ovarian cancer, liver cancer, kidney cancer, skin cancer and cancer of the vulva.